UNITED STATES PATENT AND TRADEMARK OFFICE

UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS P.O. Box 1450 Alexandria, Virginia 22313-1450 www.uspto.gov

				
APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/699,679	10/30/2000	Evan C. Unger	UNGR-1598	8248
28213 DLA PIPER U	7590 10/02/2007 S LLP	EXAMINER		
4365 EXECUTIVE DRIVE SUITE 1100			SCHLIENTZ, LEAH H	
SAN DIEGO, CA 92121-2133			ART UNIT	PAPER NUMBER
			1618	
			MAIL DATE	DELIVERY MODE
	•		10/02/2007	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

		Application No.	Applicant(s)			
Office Action Summary		09/699,679	UNGER ET AL.			
		Examiner	Art Unit			
		Leah Schlientz	1618			
	The MAILING DATE of this communication app					
Period fo						
WHIC - Exter after - If NC - Failu Any	ORTENED STATUTORY PERIOD FOR REPLY CHEVER IS LONGER, FROM THE MAILING DANS IN THE MAIL	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tim rill apply and will expire SIX (6) MONTHS from cause the application to become ABANDONEI	I. lely filed the mailing date of this communication. (35 U.S.C. § 133).			
Status	•					
1)⊠	Responsive to communication(s) filed on 25 Ju	ine 2007.				
·	This action is FINAL . 2b)⊠ This action is non-final.					
3)	•—					
	closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.					
Dispositi	on of Claims					
4)⊠	Claim(s) 3,4,12,13,17,22-35,61 and 63-81 is/ar	e pending in the application.				
4a) Of the above claim(s) <u>12 and 13</u> is/are withdrawn from consideration.						
5)□	Claim(s) is/are allowed.					
6)⊠	Claim(s) 3,4,17,22-35,61 and 63-81 is/are reject	cted.				
7)	Claim(s) is/are objected to.					
8)□	Claim(s) are subject to restriction and/or	election requirement.				
Applicati	on Papers					
9)[The specification is objected to by the Examine	r.				
	The drawing(s) filed on is/are: a)☐ acce		Examiner.			
	Applicant may not request that any objection to the	drawing(s) be held in abeyance. See	37 CFR 1.85(a).			
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).						
11)☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.						
Priority u	ınder 35 U.S.C. § 119					
12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of:						
	1. Certified copies of the priority documents have been received.					
2. Certified copies of the priority documents have been received in Application No						
3. Copies of the certified copies of the priority documents have been received in this National Stage						
application from the International Bureau (PCT Rule 17.2(a)).						
* See the attached detailed Office action for a list of the certified copies not received.						
Attachment		_				
1) Unotice of References Cited (PTO-892) 4) Interview Summary (PTO-413) Paper No(s)/Mail Date						
3) 🔲 Inform	nation Disclosure Statement(s) (PTO/SB/08)	5) 🔲 Notice of Informal Pa				
	No(s)/Mail Date	6) Other:				

DETAILED ACTION

Continued Examination Under 37 CFR 1.114

A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on 6/25/2007 has been entered.

Status of Claims

Claims 3, 4, 17 and 66 have been amended. Claims 3, 4, 12, 13, 17, 22 – 35, 61, and 63 – 81 are pending, of which claims 12 and 13 are withdrawn from consideration at this time as being drawn to a non-elected invention. Claims 3, 4, 17, 22 – 35, 61, and 63 – 81 are readable upon the elected invention and are examined herein on the merits for patentability.

The indicated allowability of claims 66 – 81 is withdrawn in view of further consideration of the prior art of record. Rejections based on the cited reference(s) follow.

Response to Arguments

Applicant's arguments, filed 6/25/2007, with respect to the rejection of claims 3, 4, 6 – 10, 17, 22 – 35, 61 and 63 – 65 under 35 USC 103(a) as being unpatentable over Unger *et al.* (WO 96/40285) have been fully considered but they are not persuasive for reasons set forth hereinbelow.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

The factual inquiries set forth in *Graham* v. *John Deere Co.*, 383 U.S. 1, 148 USPQ 459 (1966), that are applied for establishing a background for determining obviousness under 35 U.S.C. 103(a) are summarized as follows:

- 1. Determining the scope and contents of the prior art.
- 2. Ascertaining the differences between the prior art and the claims at issue.
- 3. Resolving the level of ordinary skill in the pertinent art.
- 4. Considering objective evidence present in the application indicating obviousness or nonobviousness.

Claims 3, 4, 17, 22 – 35, 61, and 63 – 81 are rejected under 35 U.S.C. 103(a) as being unpatentable over Unger (WO 96/40285).

Unger discloses novel targeted compositions which may be used for diagnostic and therapeutic use, such as for therapeutic ultrasound (see abstract). Unger teaches

that the composition can comprise a vesicle composition having an aqueous carrier, vesicles comprising a lipid and a gas, such as gas-filled liposomes (see page 4, lines 20 - 30, page 9, lines 15 - 25 and page 12, lines 25 - 33), and thus teaches a targeted vesicle composition for therapeutic or diagnostic use *in vivo* having an aqueous carrier and gas filled liposomes, as recited in claims 17 and 66. Unger further teaches that exemplary lipids that can be used to prepare the liposomes comprise phosphatidyl chlolines such as dioleylphosphatidylcholine, dimyristoylphosphatidylcholine, and others (see page 23, lines 15 - 34).

Unger further teaches that the composition comprises a compound having the formula:

Where Q is a targeting ligand (see page 61, line 1 through page 62, line 33 and claim 136). It is noted that the compound as recited in claim 136 of Unger meets the limitation of the instant structure in that the carbon atom of the structure is linked to R_2 - X_1 - R_1 at one end and X_1 - R_1 at the other, and R_3 at the third position. Unger defines R_2 as being an alkylene moiety from 1-30 carbon atoms, which encompasses the instantly claimed species of R^3 as ethylene. Regarding the moieties R^1 and R^4 as recited in instant claims 17 or 66, Unger teaches that X_1 may be $-C(=X)_5$ - X_4 , where X_5 can be O, X_4 can be $-NR_4$ - and R_4 can be a hydrogen or alkyl of 1-10 carbon atoms, as in the instantly claimed structure wherein R^2 and R^5 are hydrogen lower alkyl, and thus

Application/Control Number: 09/699,679

Art Unit: 1618

teaches the moiety -C(=O)-N(alkyl)- that meets the limitation of the fragment $C(=O)-N(R^2)$ -, as recited in claims 17 and 66. Regarding the acyl moieties R^1 and R^4 , Unger teaches that the alkyl group connected to -C(=O)-N(alkyl)- (and that thus forms an acyl group) can comprise an alkyl of 1 to 50 carbons, and thus teaches an acyl group that encompasses the claimed species of R^1 that has from 16 to 23 carbons, including 18. R_3 may be hydrogen.

Unger teaches a bond between the central carbon C and moiety M, and thus teaches R^6 is a direct bond, as instantly claimed in claims 17 and 66. Regarding moiety X^1 as recited in claims 17 and 66, Unger teaches that moiety M may be $-R_5$ -C(= X_5)- X_4 , where R_5 can be a direct bond, X_5 can be O, and X_4 can be NR₄, with R₄ being hydrogen or an alkyl from 1-10 carbon atoms. Thus Unger teaches that M can be the moiety C(=O)-N(alkyl)-, which meets the limitation of X^1 , as instantly claimed. Unger teaches that X_2 connected to M can be a direct bond, and moiety Z can be a hydrophilic polymer, which is preferably PEG (see page 62, lines 25 – 27 and claim 146).

Regarding the moieties R^7 and X^2 , as recited in claims 17 and 66, Unger teaches that a moiety X_3 is present between the hydrophilic polymer, Z, and the peptide Q. Unger teaches that moiety X_3 can be $-R_5$ - $C(=X_5)$ - X_4 , wherein R_5 is alkyl (preferably C_1 or C_2), X_5 is O, and X_4 is N, and thus teaches X_3 can be (alkyl)C(=O)-N- (pages 61 – 64), which is directly overlapping in scope with instantly claimed portion of the compound R^7 - X^2 -peptide, wherein R^7 is CH_2CH_2 , X_2 is C(=O), and N may be inherently donated from the targeting peptide via an amide bond in the instantly claimed structure.

Regarding the recitation in claims 17 and 66 that the targeting ligand T is a peptide having the sequence CRGDC, wherein the two cysteines are linked together via a disulfide linkage, Unger teaches that ligands useful for targeting the GPIIbIIa receptor include peptides flanked by cysteine residues that are capable of forming cyclic disulfides, such as cyclic, disulfide-bonded forms with the sequence Arg-Gly-Asp (i.e. RGD) (see page 57, lines 23 - 33 and page 55, lines 20 - 30), and thus teaches providing a peptide of sequence CRGDC as a targeting ligand for targeting the GPIIbIIa receptor.

Unger does not teach a specific embodiment of the compound having a combination of the specifically claimed targeting ligand that targets the GPIIbIIa receptor and the hydrophilic polymer, as in the elected species of compound.

However, it is considered that one of ordinary skill in the art at the time of the instant invention would have found it obvious to provide the compound that meets the limitation of formula (IV), as claimed and in particular, the elected species of such formula wherein the specific targeting ligand that targets the GPIIbIIa receptor and the hydrophilic polymer that is polyethylene glycol, because Unger teaches the compound having the structure that overlaps with and/or meets the limitations of the instant claims 17 and 66, and furthermore teaches that the compound can comprise targeting ligands that include the GPIIbIIa receptor as claimed and the polyethylene glycol hydrophilic polymer as claimed, and teaches such compounds are useful in a vesicle composition for diagnostic and therapeutic use. Accordingly, it is considered that one of ordinary skill in the art would have been motivated to provide the claimed compound with the

expectation of providing a suitable compound for formulation in a vesicle composition for diagnostic use. Accordingly, claims 17 and 66 are obvious over the teachings of Unger et al.

Regarding claims 3, 4, 61 and 63, Unger teaches the compositions of the claims insofar as they read on the elected species, as discussed above.

Regarding claim 22 and 68, Unger teaches that the phosphatidylcholine provided in the composition can be dipalmitoylphosphatidylcholine (page 23, lines 15-34). Regarding claims 23 – 24 and 69 – 70. Unger teaches that the lipids can comprise dipalmitovlphosphatidylethanolamine (page 23, lines 15-33). Regarding claims 25 and 71, Unger teaches that the lipids can comprise dipalmitoylphosphatidic acid (page 23, lines 15-34).

Regarding claims 26-29 and 72 – 75, Unger teaches that the vesicles can comprise a gas such as perfluorocarbon, including perfluoromethane, perfluoropropane, perfluorobutane, etc. (page 32, lines 3 –18). Regarding claims 30-33 and 76 – 79, Unger teaches the gas can be derived from a gaseous precursor such as perfluoropentane that is converted to a gas at 37 C (page 33, lines 16-33). The composition can also comprise bioactive agents such as urokinase, heparain, etc. (page 83, lines 9-24).

Regarding claims 64-65, the targeting ligand can comprise a peptide having the claimed number of amino acids, and can be cyclized (page 55, line 1 – page 59, line 20).

Application/Control Number: 09/699,679

Art Unit: 1618

Applicant argues, on page 12 of the Response that claim 17 as amended, now recites a general structure where "X is C(=0);" " R^7 is (CH_2CH_2) ;" and "P is PEG;" and that Unger fails to describe or suggest a composition that comprises a compound that satisfies all these limitations. Applicant contends that with respect to moiety X^2 , all that Unger teaches is the moiety C(=0)-N-alkyl, and with respect to the polymeric moiety, all that Unger teach is generic PEG.

This is non-persuasive because, regarding instantly claimed moiety X^2 , Unger teaches that moiety X_3 can be $-R_5$ - $C(=X_5)$ - X_4 , wherein R_5 is alkyl, X_5 is O, and X_4 is N, and thus teaches X_3 can be $CH_2CH_2C(=O)$ -N- (pages 61 – 64), which is directly overlapping in scope with instantly claimed portion of the compound R^7 - X^2 -peptide, wherein R^7 is CH_2CH_2 , X_2 is C(=O), and N may be inherently donated from the targeting peptide via an amide bond in the instantly claimed structure, as set forth above. Regarding PEG, all that is claimed with respect to moiety P is generic PEG.

Conclusion

No claims are allowed at this time.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Leah Schlientz whose telephone number is 571-272-9928. The examiner can normally be reached on Monday - Friday 8 AM - 5 PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Michael Hartley can be reached on 571-272-0616. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Application/Control Number: 09/699,679 Page 9

Art Unit: 1618

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

LHS

MICHAEL G. HARTLEY
SUPERVISORY PATENT EXAMINER